

Food and Drug Administration Rockville MD 20857

Re: Requip

Docket No.: 98E-0478

DEC - 4 1998

The Honorable Bruce Lehman
Assistant Secretary of Commerce and
Commissioner of Patents and Trademarks
Box Pat. Ext.
Assistant Commissioner for Patents
Washington, DC 20231

Dear Commissioner Lehman:

ASSISTABLE SECRETARY

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TO ADEMARK OFFICE

This is in regard to the application for patent term extension for U.S. Patent No. 4,452,808, filed by SmithKline Beecham Corp., under 35 U.S.C. § 156 et seq. We have reviewed the dates contained in the application and have determined the regulatory review period for Requip, the human drug product claimed by the patent.

The total length of the regulatory review period for Requip is 3,356 days. Of this time, 2,729 days occurred during the testing phase and 627 days occurred during the approval phase. These periods of time were derived from the following dates:

1. The date an exemption under subsection 505(i) of the Federal Food, Drug, and Cosmetic Act involving this drug product became effective: July 14, 1988.

The applicant claims July 10, 1988, as the date the Investigational New Drug application (IND) became effective. However, FDA records indicate that the IND effective date was July 14, 1988, which was thirty days after FDA receipt of the IND.

2. The date the application was initially submitted with respect to the human drug product under section 505 of the Federal Food, Drug, and Cosmetic Act: January 2, 1996.

FDA has verified the applicant's claim that the New Drug Application (NDA) for Requip (NDA 20-658) was initially submitted on January 2, 1996.

3. The date the application was approved: September 19, 1997.

FDA has verified the applicant's claim that NDA 20-658 was approved on September 19, 1997.

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This determination of the regulatory review period by FDA does not take into account the effective date of the patent, nor does it exclude one-half of the testing phase as required by 35 U.S.C. § 156(c)(2).

Please let me know if we can be of further assistance.

Sincerely yours,

Thomas J. McGinnis, R.Ph.

Deputy Associate Commissioner

for Health Affairs

cc: Stephen Venetianer

SmithKline Beecham Corp

Corp. Intellectual Property UW2220

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